

## **Instructions for Completion of the Patient Safety Annual Facility Survey for LTAC (CDC 57.150)**

Data Field	Instructions for Form Completion
Facility ID #	Required. The NHSN-assigned facility ID will be auto-entered by the computer.
Survey Year	<i>Required.</i> Select the calendar year for which this survey was completed. The survey year should represent the last full calendar year. For example, in 2014, a facility would complete a 2013 survey.
Facility Characteristics	
Ownership (check one)	<ul> <li>Required. Select the appropriate ownership of this facility:</li> <li>For profit</li> <li>Not for profit, including church</li> <li>Government</li> <li>Veterans Affairs</li> </ul>
Affiliation (check one)	<ul> <li>Required. Select the appropriate affiliation for this facility:</li> <li>Independent – The facility is a stand-alone facility that does not share a building, staff, or policies (such as infection control) with any other healthcare institution.</li> <li>Hospital system – The facility is affiliated with a local healthcare system. Facility shares policies (such as infection control) with other institutions within the hospital system. Facility may or may not share staff as well as a building with other facilities that are part of that hospital system.</li> <li>Multi-facility organization (specialty network) – The facility is part of a regional or national network of specialty facilities. Facilities share policies (such as infection control), corporate leadership, and a common business structure.</li> </ul>
Setting/Classification:	Required. Select the physical setting of the facility: free-standing or within a hospital.
If classified as "Free-standing", does your LTAC hospital share physical housing with one or more of the following on-site facilities or units? (check all that apply)	Conditionally Required. If facility is classified as free-standing, select one or more of the following facility or unit types that share physical housing with your LTAC:  No (none) Skilled nursing facility (SNF)/nursing home Residential facility (assisted living) Inpatient rehabilitation facility Neuro-behavioral unit or facility Other: specify



Facility Characteristics (continued)		
If classified as "Within a	Conditionally Required. If facility is classified as within a hospital, indicate	
hospital", is your LTAC hospital	'Yes' or 'No' if it is:	
located:	• In a building that does not provide acute care services (e.g., psychiatric	
	hospital)	
	Near (but not within) an acute care hospital	
	Note: These questions are to clarify the physical set-up of each HOSP-LTAC.	
	All separately licensed LTAC hospitals should be enrolled within NHSN as a	
	separate LTAC facility type regardless of the physical setting of the facility.	
Number of Patient Days	Required. Enter the total number of patient days for your hospital during the	
	last full calendar year.	
Number of Admissions	Required. Enter the total number of inpatient admissions for your hospital	
	during the last full calendar year.	
Average daily census	Required. Enter the average number of patients housed each day during the last full calendar year. Please round to the nearest whole number.	
Numbers of LTAC beds in the	Required. Enter the total number of LTAC beds in each on the following	
following categories (categories	categories during the last full calendar year:	
should equal total number of	• Intensive care unit (ICU) or critical care beds	
beds)	High observation/special care/high acuity beds (not ICU)	
	Other LTAC beds	
Total number of LTAC beds	Required. The total number of LTAC beds in the facility during the last full	
(licensed capacity)	calendar year will be automatically summed based on the above counts.	
Number of single occupancy	Required. Enter the total number of single occupancy rooms during the last	
rooms	full calendar year.	
Number of admissions with the	Required. Enter the number of admissions with the <b>primary diagnosis</b> for	
primary diagnosis for each	each of the following categories:	
category	Ventilator dependence     Home distance	
	<ul><li>Hemodialysis</li><li>Chronic wound care</li></ul>	
Fooility Miorobists of Late	• Long term IV therapy	
	ory Practices. Completion of this section requires the assistance from the	
microbiology laboratory. Question majority of the last full calendar	ons should be answered based on the testing methods that were used for the	
	Required. Select 'Yes' if your laboratory performs antimicrobial susceptibility	
1. Does your facility have its own laboratory that	testing; otherwise, select 'No'.	
performs antimicrobial	lesting, otherwise, select two.	
susceptibility testing?		
susceptionity testing:		
If No, where is the facility's	Conditionally Required. If 'No', select the location where your facility's	
antimicrobial susceptibility	antimicrobial susceptibility testing is performed: Affiliated medical center,	
testing performed? (check	Commercial referral laboratory, or Other local/regional, non-affiliated	
one)	reference laboratory. If multiple laboratories are used indicate the laboratory	
,	which performs the majority of the bacterial susceptibility testing. You must	
	complete the remainder of this survey with assistance from your outside	
	laboratory.	



Fac	cility Microbiology Laborato	ory Practices (continued)
	Does the laboratory use CLSI (formerly NCCLS) antimicrobial susceptibility standards? If Yes, specify which version of the M100 document the laboratory uses.	Required. Select 'Yes' if your laboratory uses CLSI antimicrobial susceptibility standards; otherwise, select 'No'.  Conditionally Required. If 'Yes', specify the version used by your laboratory or the referral laboratory.
3.	For the following organisms please indicate which methods are used for (1) primary susceptibility	Required. Select from the choices listed the appropriate (1) primary susceptibility testing and (2) secondary, supplemental, or confirmatory testing method (if performed) for each organism.
	testing and (2) secondary, supplemental, or confirmatory testing (if performed)	Note: Repeat tests using the primary method should not be indicated as secondary methods; instead indicate in the 'Comments' column the number of times repeat testing is done using the same primary method.
		If your laboratory does not perform susceptibility testing, please indicate the methods used at the referral laboratory. If 'Other' is selected as the method for any pathogen, use the 'Comments' column to describe the method used.
4.	Has your laboratory implemented the revised cephalosporin and monobactam breakpoints for Enterobacteriaceae recommended by CLSI as of 2010?	
5.	Has your laboratory implemented the revised	Required. Select 'Yes' if your laboratory has implemented the revised carbapenem breakpoints for Enterobacteriaceae recommended by CLSI as of 2010; otherwise, select 'No'.
6.	Does your laboratory perform a special test for carbapenemase production?	Required. Select 'Yes' if your laboratory performs a special test for carbapenemase production; otherwise, select 'No'.
		Conditionally Required. If 'Yes', specify what is done if carbapenemase production is detected.
	If Yes, which test is routinely performed to detect carbapenemase (check all that apply)?	Conditionally Required. If 'Yes', specify which test is performed to detect carbapenemase.



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Fac	acility Microbiology Laboratory Practices (continued)		
7.	Does your laboratory perform colistin or polymyxin B susceptibility testing for drug-resistant gram negative bacilli? If Yes, indicate methods (check all that apply).	Required. Select 'Yes' if your laboratory performs colistin or polymyxin B susceptibility testing for drug-resistant gram negative bacilli; otherwise, select 'No'.  Conditionally Required. If 'Yes', select the method(s) used from the choices provided. If 'Other' is selected, please specify.	
8.	Does your facility have its own laboratory that performs antifungal susceptibility testing for <i>Candida</i> species? If No, where your facility's antifungal susceptibility testing is performed? (check one).	Required. Select 'Yes' if your laboratory performs antifungal susceptibility testing for Candida species; otherwise, select 'No'.  Conditionally Required. If 'No", select one of the choices provided.	
9.	If antifungal susceptibility testing is performed at your facility or an outside laboratory, what methods are used? (check all that apply)	Conditionally Required. Select from the choices listed the method(s) of antifungal susceptibility testing performed at your facility or an outside laboratory. If 'Other' is selected, please specify.	
	Candida spp. cultured from normally sterile body sites (such as blood), without needing a specific order or request for susceptibility testing from the clinician? If Yes, what antifungal drugs are tested automatically/reflexively? (check all that apply)	Required. Select 'Yes' if your laboratory or outside laboratory automatically/reflexively performs antifungal susceptibility testing for Candida species which are from normally sterile body sites (such as blood), without needing a specific order or request for susceptibility testing from the clinician; otherwise, select 'No'.  Conditionally Required. If 'Yes', select from the choices listed the antifungal drugs which are tested automatically/reflexively. If 'Other' is selected, please specify.	
11.	What is the primary testing method for <i>C. difficile</i> used most often by your facility's laboratory or the outside laboratory where your facility's testing is performed? (check one)	Required. Select from the choices listed the testing methods used to perform <i>C. difficile</i> testing by your facility's laboratory or the outside laboratory where your facility's testing is done. If 'Other' is selected, please specify.  Note: "Other" should not be used to name specific laboratories, reference laboratories, or the brand names of <i>C. difficile</i> tests; most methods can be categorized accurately by selecting from the options provided. Please ask your laboratory or conduct a search for further guidance on selecting the correct option to report.	



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Facility Microbiology Laborate	
12. Does your facility produce an antibiogram (i.e., cumulative antimicrobial susceptibility report)?	Required. Select 'Yes' if your facility produces an antibiogram; otherwise select 'No'.
If Yes, is the antibiogram produced at least annually?	Conditionally Required. If 'Yes', indicate whether the antibiogram is produced at least annually.
If Yes, are data stratified by hospital location?  If No, please identify any obstacle(s) to producing an	Conditionally Required. If 'Yes', indicate whether antibiogram data are stratified by hospital location.  Conditionally Required. If 'No', indicate the obstacle(s) to producing an antibiogram at your facility. If 'Other' is selected, please specify.
antibiogram. (Check all that apply)	
	impletion of this section may require assistance from the Infection Preventionist,
	nfection control personnel, and/or Quality Improvement Coordinator. Questions epolicies and practices that were in place for the majority of the last full calendar
13. Number of infection preventionists (IPs) in facility	Required. Enter the number of individuals (full-time employees) who work in the infection prevention department of the hospital as infection prevention professionals. Certification in infection control, the CIC credential, is not required to be considered an "IP" on this survey.
a. Total hours per week performing surveillance	Enter the number of hours per week engaged in activities designed to find and report healthcare-associated infections (in the hospital) and the appropriate denominators. Total should include time to analyze data and disseminate results.
b. Total hours per week for infection control activities other than surveillance	Enter the number of hours per week spent on infection prevention and control activities other than surveillance. These activities include, but are not limited to, education, prevention, meetings, etc.
14. Does your facility perform active surveillance testing	<i>Required.</i> Select from the choices listed, all MDRO(s) for which newly-admitted patients are tested for colonization:
(culturing) of new patients	Methicillin-resistant Staphylococcus aureus (MRSA)
on admission for	• Vancomycin-resistant Enterococcus (VRE)
colonization with any of the	··· <del>··</del> · · · · · · · · · · · · · · · ·
following multidrug- resistant organisms	Other multidrug-resistant gram-negative rods
(MDROs)? (check all that apply)	We do not screen new admissions for MDROs
For detailed description about th	ne use of Contact Precautions, please refer to the CDC/HICPAC 2007 Guideline
for Isolation Precautions: Preve (http://www.cdc.gov/hicpac/pdf/v	nting Transmission of Infectious Agents in Healthcare Settings solation/Isolation2007.pdf).
15. Does the facility routinely place patients infected or colonized with MRSA in contact precautions? (check	Required. Select 'No' if your facility does not routinely place any patient infected or colonized with MRSA in Contact Precautions; otherwise, select the single best choice from the choices listed that most accurately describes the primary indication for placing patients with MRSA on Contact Precautions at your facility.
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Inf	Infection Control Practices (continued)		
16.	Does the facility routinely place patients infected or colonized with VRE in contact precautions? (check one)	Required. Select 'No' if your facility does not routinely place any patient infected or colonized with VRE in Contact Precautions; otherwise, select the single best choice from the choices listed that most accurately describes the primary indication for placing patients with VRE on Contact Precautions at your facility.	
17.	Does the facility routinely place patients infected or colonized with CRE in contact precautions? (check one)	Required. Select 'No' if your facility does not routinely place any patient infected or colonized with CRE in Contact Precautions; otherwise, select the single best choice from the choices listed that most accurately describes the primary indication for placing patients with CRE on Contact Precautions at your facility.	
18.	Does the facility routinely place patients infected or colonized with ESBL-producing or extended spectrum cephalosporin resistant Enterobacteriaceae in contact precautions? (check one)	Required. Select 'No' if your facility does not routinely place any patient infected or colonized with ESBL-producing or extended spectrum cephalosporin-resistant Enterobacteriaceae in Contact Precautions; otherwise, select the single best choice from the choices listed that most accurately describes the primary indication for placing patients with ESBL-producing or extended spectrum cephalosporin-resistant Enterobacteriaceae on Contact Precautions at your facility.	
19.	Does the facility routinely perform screening cultures for CRE?	Required. Select 'Yes' if your facility <u>routinely</u> obtains screening cultures from patients for CRE (e.g., rectal, perirectal, or stool cultures); otherwise, select 'No.'	
	If Yes, in which situations does the facility routinely perform screening cultures for CRE? (check all that apply)	Conditionally Required. If 'Yes', select <u>all</u> the situations in which your facility would <u>routinely</u> obtain screening cultures from patients for CRE. If 'Other' is selected, please specify the situation(s) in which CRE screening is performed.  Note: 'Epidemiologically-linked' patients refer to contacts of the patient with newly identified CRE. This might include current or prior roommates or patients who shared the same healthcare personnel or patients who are located on the same unit or ward.	
20.	Does the facility use chlorhexidine bathing on any patient to prevent transmission of MDROs in your hospital?	Required. Select 'Yes' if your facility <u>routinely</u> uses chlorhexidine bathing on any patient in any ward or unit as an intervention to prevent the transmission of any MDRO; otherwise, select 'No'.	
21.	Are results rapidly communicated (generally within 4 hours) to infection prevention staff and/or clinical staff when MDROs are identified from clinical or screening cultures in the	Required. Select 'Yes' if the laboratory that performs clinical culture testing for your facility routinely notifies relevant staff (either the Infection Prevention staff and/or clinical staff) in a timely manner (e.g., within 4 hours) when an MDRO is identified from clinical or surveillance cultures; otherwise, select 'No.'	
	laboratory? If Yes, for which MDROs? (check all that apply)	Conditionally required. If 'Yes', select all the MDROs for which timely notification of relevant staff is performed by the laboratory. If 'Other' is selected, please specify the MDRO(s) for which this would apply.	



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Inf	Infection Control Practices (continued)		
22.	When a patient with an MDRO is transferred to another facility, does your facility communicate the patient's MDRO status to the receiving facility at the time of transfer?	Required. Select 'Yes' if your facility <u>routinely</u> communicates the MDRO status of a patient known to be colonized or infected with an MDRO to the receiving facility at the time of patient transfer; otherwise, select 'No'.	
23.	Among patients with an MDRO admitted to your facility from another healthcare facility, what percentage of the time does the facility receive information from the transferring facility about the patient's MDRO status?	Required. Enter the estimated percentage of the time that your facility receives information from a transferring facility about the MDRO status of a patient known to be colonized or infected with an MDRO.	
An		es. Completion of this by section may require assistance from the pharmacy and/or	
to o Ster sho yea	optimize and measure antibiotowardship Programs ( <a href="http://www.uld">http://www.uld</a> be answered based on the r.	mittee. Antibiotic Stewardship refers to a coordinated, multidisciplinary approach ic use. For further information, refer to Core Elements of Hospital Antibiotic <a href="www.cdc.gov/getsmart/healthcare/implementation/core-elements.html">www.cdc.gov/getsmart/healthcare/implementation/core-elements.html</a> ). Questions policies and practices that were in place for the majority of the last full calendar	
24.	Does your facility have a written statement of support from leadership that supports efforts to improve antibiotic use (antibiotic stewardship)?	Required. Select 'Yes' if there is written evidence of senior-level management support focused on antibiotic use prescribing (e.g., formal letter of support for efforts to improve antibiotic use, written communication to hospital staff that encourages optimal antibiotic prescribing, communication of support that reaches staff beyond those who receive executive-level meeting notes); otherwise, select 'No'.	
25.	Is there a leader responsible	Required. Select 'Yes' if any individual has been identified as a lead to antibiotic stewardship activities as evidenced by responsibility for improving antibiotic use in the job description or performance review, authority to coordinate activities of staff from multiple departments (e.g. laboratory, pharmacy, information technology), and/or responsibility to report to senior level management on program planning and outcomes.	
	If Yes, what is the position of this leader? (check one)	Conditionally Required. If 'Yes', specify the qualification or job title of the leader(s). More than choice one may be selected. If 'Other' is selected, please specify the position.	
26.	Is there at least one pharmacist responsible for improving antibiotic use at your facility?	Required. Select 'Yes' if your facility has at least one pharmacist who dedicates time distinct from general pharmacy duties to educate staff, and track or monitor antibiotic use to ensure optimal prescribing practices; otherwise, select 'No'.	



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Ant	tibiotic Stewardship Practic	es (continued)
	stewardship activities?	Required. Select 'Yes' if any individual was given salary support at least 4 hours per week (0.1 full-time employees) to engage in duties to improve or monitor antibiotic use that are not part of their general clinical duties; otherwise, select 'No'.
28.	Does your facility have a policy that requires prescribers to document an indication for all antibiotics in the medical record or during order entry?	Required. Select 'Yes' if your facility has a policy requiring documentation of dose, duration and indication for all antibiotics in the medical record or during order entry; otherwise, select 'No'.
	If Yes, has adherence to the policy to document an indication been monitored?	Conditionally Required. If 'Yes' to question 28, select 'Yes' if charts have been audited to confirm documentation of dose, duration, and indication in patient medical records; otherwise, select 'No'.
29.		Required. Select 'Yes' if there are facility-specific recommendations for antibiotic treatment selection based on national guidelines and local susceptibility reports for ANY common clinical conditions (e.g., community required pneumonia, urinary tract infections, or skin and soft tissue infections); otherwise, select 'No'.
	If Yes, has adherence to facility-specific treatment recommendations been monitored?	Conditionally Required. If 'Yes' to question 29, select 'Yes' charts have been audited to confirm adherence to facility-specific treatment guidelines for ANY of the common clinical conditions listed above; otherwise, select 'No'.
30.	Is there a formal procedure for all clinicians to review the appropriateness of all antibiotics at or after 48 hours from the initial orders (e.g. antibiotic time out)?	Required. Select 'Yes' if your facility has developed a standardized way for clinicians on the treating team (or attending physician? or physician of record?) to reassess the continuing need and choice of antibiotics at or after 48 hours after the initial orders (to confirm indication, review microbiology results, and review antibiotic choice, dose, and duration); otherwise, select 'No'.
31.	agents need to be approved	Required. Select 'Yes' if your facility has at least one antibiotic agent that requires a physician or pharmacist to review and approve administration of the drug due to its spectrum of activity, cost, or associated toxicities; otherwise, select 'No'.
32.	Does a physician or pharmacist review courses of therapy for specified antibiotic agents and communicate results with prescribers (i.e., audit with feedback) at your facility?	Required. Select 'Yes' if your facility had physicians or pharmacists knowledgeable in antibiotic use, and not part of the treating team, review courses of therapy for specified antibiotic agents and communicate the results to prescribers (such as audit with feedback); otherwise, select 'No'.



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An	Antibiotic Stewardship Practices (continued)		
33.		Required. Select 'Yes' if your facility monitors antibiotic use or consumption at the unit, service, and/or facility wide level at least quarterly; otherwise, select 'No'.	
		Conditionally Required. If 'Yes', select from the choices of listed antibiotic use metrics. Days of Therapy (also known as Antimicrobial Days) is defined by any amount of a specific antimicrobial agent administered in a calendar day to a particular patient (i.e., each antimicrobial agent administered to a patient counted as one day of therapy). The Defined Daily Dose is the assumed average maintenance dose per day for a drug used for its main indication in adults and is derived from the total number of grams of each antibiotic purchased, dispensed, or administered. If 'Other' is selected, please specify the method(s) or metric(s) used.	
	If Yes, are facility- and/or unit-specific reports on antibiotic use shared with prescribers?	Conditionally Required. Select 'Yes' if facility and/or unit-specific reports on antibiotic use are shared with prescribers (individually, by service line, by medical group, etc.); otherwise, select 'No'.	
34.	feedback by the stewardship program about how they can	Required. Select 'Yes' if prescribers (individually, by service line, by medical group, etc.) have received feedback based on observed practices through inperson, telephone, written or electronic communication about how they can improve their antibiotic prescribing; otherwise, select 'No'.	
35.	program provided education	Required. Select 'Yes' if your facility stewardship program has provided education on how to improve antibiotic use to clinicians and other relevant staff (e.g. Grand Rounds, in-service training, or direct instruction); otherwise, select 'No'.	